## **Amendments To The Claims:**

This listing of claims will replace all prior versions, and listings, of claims in the application:

## In the Claims:

- 1. (Currently amended) A solid pharmaceutical composition for oral administration comprising (2S)-2-amino-4-{[2-(ethanimidoylamino)ethyl]thio} butanoic acid, a pharmaceutically acceptable bulking agent and one or more antioxidants or chelating agents, wherein the (2S)-2-amino-4-{[2-(ethanimidoylamino)ethyl]thio}butanoic acid is in the form of its (1:1) compound with phosphoric acid, or a solvate thereof.
- 2. (Canceled).
- 3. (Previously presented) The pharmaceutical composition as claimed in claim 2 wherein the solvate is a hydrate.
- 4. (Previously presented) The pharmaceutical composition as claimed in claim 3 wherein the hydrate is the monohydrate.
- 5. (Previously presented) The pharmaceutical composition as claimed in claim 3 wherein the hydrate is the trihydrate.
- 6. (Previously presented) The pharmaceutical composition as claimed in claim 1 wherein the (2S)-2-amino-4-{[2-(ethanimidoylamino)ethyl]thio}butanoic acid comprises from about 0.1 to about 5% by weight, the pharmaceutically acceptable bulking agent comprises from about 80 to about 99.5% by weight, and the antioxidant, chelating agent, or mixture thereof comprises from about 0.005 to about 5% by weight, based on the dry weight.
- 7. (Previously presented) The pharmaceutical composition as claimed in claim 1 wherein the antioxidants or chelating agents are selected from the group comprising EDTA, malic acid, ascorbic acid and mixtures thereof.

- 8. (Previously presented) The pharmaceutical composition as claimed in claim 1 wherein the pharmaceutically acceptable bulking agent comprises microcrystalline cellulose, starch or a mixture thereof.
- 9. (Previously presented) A method for the treatment of a clinical condition in a mammal, for which an inhibitor of nitric oxide synthase is indicated, which comprises administration of a pharmaceutical composition as claimed in claim 1.
- 10. (Previously presented) The method as claimed in claim 9 wherein the clinical condition is selected from the group consisting of arthritis, asthma, rhinitis, chronic obstructive pulmonary disease, ileus, migraine, pain and irritable bowel syndrome.
- 11. (Cancelled)
- 12. (Cancelled)
- 13. (Cancelled)
- 14. (Cancelled)
- 15. (Cancelled)
- 16. (Previously presented) The method as claimed in claim 9 wherein said mammal is a human.
- 17. (Canceled).